

From the
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

PCT

To:

GERVASI, Gemma
NOTARBARTOLO & GERVASI S.P.A.
Corso di Porta Vittoria, 9
I-20122 Milano
ITALIE

NOTARBARTOLO & GERVASI
MILANO
R E C E I V E D

24 GEN. 2005

NOTIFICATION OF TRANSMITTAL OF
THE INTERNATIONAL PRELIMINARY
EXAMINATION REPORT

(PCT Rule 71.1)

Date of mailing
(day/month/year)

21.01.2005

Applicant's or agent's file reference
3585PTWO/AG/la

IMPORTANT NOTIFICATION

International application No.
PCT/EP 03/11642

International filing date (day/month/year)
21.10.2003

Priority date (day/month/year)
21.10.2002

Applicant

L. MOLteni & C. DEI FRATELLI ALITTI SOCIETA DI...

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.

4. REMINDER

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices) (Article 39(1)) (see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

The applicant's attention is drawn to Article 33(5), which provides that the criteria of novelty, inventive step and industrial applicability described in Article 33(2) to (4) merely serve the purposes of international preliminary examination and that "any Contracting State may apply additional or different criteria for the purposes of deciding whether, in that State, the claimed inventions is patentable or not" (see also Article 27(5)). Such additional criteria may relate, for example, to exemptions from patentability, requirements for enabling disclosure, clarity and support for the claims.

Name and mailing address of the international
preliminary examining authority:



European Patent Office
D-80298 Munich
Tel. +49 89 2399 - 0 Tx: 523656 epmu d
Fax: +49 89 2399 - 4465

Authorized Officer

Ullrich, J

Tel. +49 89 2399-8048



PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT (PCT Article 36 and Rule 70)

Applicant's or agent's file reference 3585PTWO/AG/1a	FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/PEA/416)	
International application No. PCT/EP 03/11642	International filing date (<i>day/month/year</i>) 21.10.2003	Priority date (<i>day/month/year</i>) 21.10.2002
International Patent Classification (IPC) or both national classification and IPC C07D487/22		
Applicant L. MOLteni & C. DEI FRATELLI ALITTI SOCIETA DI...		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 5 sheets, including this cover sheet.

☒ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of 15 sheets.

3. This report contains indications relating to the following items:

- I ☒ Basis of the opinion
- II ☐ Priority
- III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Rule 66.2(a)(ii) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain documents cited
- VII ☐ Certain defects in the international application
- VIII ☐ Certain observations on the international application

Date of submission of the demand 21.05.2004	Date of completion of this report 21.01.2005
Name and mailing address of the international preliminary examining authority: <div style="display: flex; align-items: center;"> <div> European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465 </div> </div>	Authorized Officer Boletti-Cremers, K Telephone No. +49 89 2399-8541

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/EP 03/11642

I. Basis of the report

1. With regard to the **elements** of the international application (*Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)*):

Description, Pages

1, 3, 4, 6-38 as originally filed
2, 2a, 5, 5a received on 22.11.2004 with letter of 16.11.2004

Claims, Numbers

1-21 received on 22.11.2004 with letter of 16.11.2004

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)).

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/EP 03/11642

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

☐ the entire international application,

☒ claims Nos. 17,19-21

because:

☒ the said international application, or the said claims Nos. 17,19-21 relate to the following subject matter which does not require an international preliminary examination (specify):

see separate sheet

☐ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (*specify*):

☐ the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.

☐ no international search report has been established for the said claims Nos.

2. A meaningful international preliminary examination cannot be carried out due to the failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions:

☐ the written form has not been furnished or does not comply with the Standard.

☐ the computer readable form has not been furnished or does not comply with the Standard.

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes: Claims	1-21
	No: Claims	
Inventive step (IS)	Yes: Claims	1-21
	No: Claims	
Industrial applicability (IA)	Yes: Claims	1-16,18
	No: Claims	

2. Citations and explanations

see separate sheet

POINT III.

For the assessment of the presently worded claims 17 and 19-21 on the question whether they are industrially applicable, no unified criteria exist in the PCT. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognise as industrially applicable claims to the use of a compound in medical treatment, but will allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a new medical treatment.

POINT V.

The following documents , quoted in the I.S.R., have been considered as relevant for the examination of the present application . Their numbering will be adhered to for the rest of the procedure.

- (1) WO-A-01/96343, cited in the application.
- (2) EP-A-0906 758, cited in the application .
- (3) WO-A-02/10173, cited in the application.
- (4) WO-A-98/33503.
- (5) J.A.C.S. , 114(7), 2664-9, 1992.

1. Novelty.

- 1. The photosensitiser disclosed in (1)-(4) do not fall within the scope of present claims which can therefore be regarded as novel with respect to their contents.
- 2. The same conclusion can now be drawn from the content of (5) because the 2 single compounds disclosed in (5) , namely compounds 6 and 7 of (5), have been properly disclaimed from the claimed matter on file which can now be regarded as novel with respect to the content of (5).

2. Inventiveness

In view of the comparative argumentation submitted on 16.11.2004 by the Applicant , present

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/EP 03/11642

claims can be regarded as inventive with respect to the most relevant prior art (1) and (3). (5) is to be set aside because it relates to a quite different problem and according to the EP-regional practice, such a prior art can be avoided by means of a disclaimer (as on file).

It should be reminded that present claims have not been fully searched and that an additional search, focusing on the M and the respective R and R₁ definitions will be performed in the regional proceedings to come.

JC13 Rec'd T/PTO 21 APR 2005

markedly limited efficiency and poor selectivity toward the eukaryotic cells and/or micro-organisms, and because of the prolonged persistence in the skin, which often may cause phenomena of generalised photosensitivity (Jori G., *J. Photochem. Photobiol., B: Biol.*, Vol. 36, pp. 87-93, 1996).

- 5 Thus it is evident how important it is to develop novel porphyrin compounds suitable for the use as therapeutic agents in PDT and as diagnostic agents, but not showing the limitations illustrated above.

Some substituted porphyrins and metalloporphyrins are disclosed in Dick D.L. et al. *J. Am. Chem. Soc.* 114 (1992) 2664-2669. Their encapsulation within the
10 hydrophobic cavity of cyclodextrins is reported, providing inclusion complexes capable of mimicking the activities of heme-containing proteins.

Porphyrin derivatives bearing cationic groups have been previously described (Merchat et al. *J. Photochem. Photobiol.* 32, 153-157, 1996; Merchat et al. *J. Photochem. Photobiol.* 35, 149-157, 1996) and assessed for their photodynamic
15 properties in the bacteria photoinactivation. These compounds bear trimethyl-anilinium groups or quaternary ammonium pyridinium groups in the meso-positions and therefore are endowed by a hydrophilic nature.

Other photosensitisers such as phthalocyanines having hydrophilic and/or amphiphilic characteristics are known; for example, the International Applications
20 No. WO 01/96343 and WO 02/090361, and in the US Patent No. 5,965,598, all in the name of the Applicant, disclose various evenly substituted hydrophilic phthalocyanines, as well as non centrosymmetrical phthalocyanines bearing cationic or protonable group on the macrocycle.

Summary of the invention

- 25 The Applicant has now found a novel series of photosensitizers having particularly advantageous properties compared to the known compounds.

These novel compounds have shown optimum physical-chemical features for therapeutic applications, particularly in relation to their absorption in the region of the visible spectrum, high molar extinction coefficients, high quantum yield in
30 singlet oxygen production, that is expressed by the photoinactivation of eukaryotic and prokaryotic cells.

The photosensitizers described by this invention are able to produce singlet oxygen by using various light sources and wavelengths. In particular they can be activated by visible red light radiation when the treatment of deep seated tumours on infections is required as well as by blue visible radiation or white light radiations
s when is preferable to treat by means of the photodynamic process more superficial

P is selected from the group consisting of O, CH₂, CO₂, NHCONH and CONH;

I is an integer comprised between 0 and 5;

W is selected from the group consisting of O, CO₂, CONH and NHCONH;

f is selected from between 0 and 1;

5 J is H or an alkyl group (CH₂)_q-CH₃, wherein q is an integer comprised between 0 and 20;

R₂ and R₃, equal or different from each other, are selected from between R and R₁, wherein R and R₁ are defined as above,

M is chosen from 2H and a metal selected from the group consisting of Zn, Mg, Pt, Pd, Si(OR₇)₂, Ge(OR₇)₂ and AlOR₇, wherein R₇ is chosen from between H and C1-

10 C15 alkyl,

and pharmaceutically acceptable salts thereof,

with the exception of the following compounds:

a) compound of formula (I) wherein M is 2H, R₁ = R₃ = H, R = R₂ is a group of formula (II) in which s is 1, X is O, Y is (CH₂)₃, v is 1, Z is N, n = d = 1, m is 0, and

15 R₄ = R₅ = H; and

b) compound of formula (I) wherein M is 2H, R₁ = R₃ = H, R = R₂ is a group of formula (II) in which s is 1, X is O, Y is (CH₂)₃, v is 1, Z is N, n = d = 1, m is 0, R₄ and R₅ form with Z a phthalimido group.

Further subject of the present invention are the processes for the preparation of
20 the above said compounds of formula (I), the novel intermediates in these processes and the conjugates in which the compounds of formula (I) are site specifically conjugated with bio-organic carriers, such as aminoacids, polypeptides, proteins and polysaccharides.

The present compounds of formula (I), as well as the corresponding conjugates,
25 are useful for the treatment of microbial infections (bacterial, fungal and viral), in the photodynamic treatment of tumour, pre-cancerous pathologies, and other hyperproliferative diseases.

The present compounds (I) and the corresponding conjugates are useful as well, as
30 diagnostic agents for the identification of pathologically affected areas and for photodynamic sterilization of blood and blood derivatives.

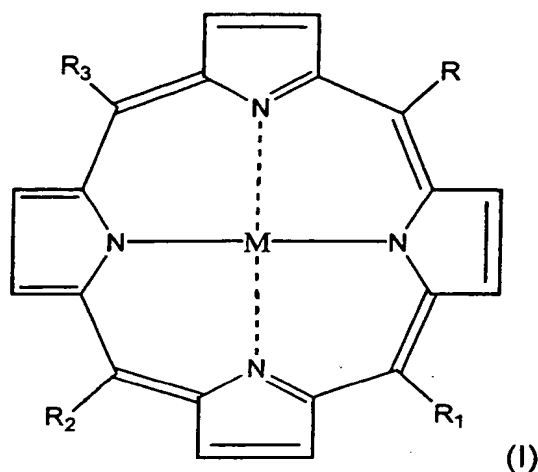
Features and advantages of the present compounds of formula (I) will be illustrated in details in the following description.

Detailed description of the invention

By "saturated or unsaturated heterocycle possibly substituted" according to the invention, an heterocycle is preferably meant, which is selected from the group consisting of morpholine, piperidine, pyridine, pyrimidine, piperazine, pyrrolidine, pyrroline, imidazole, aniline and julolidine (2,3,6,7-tetrahydro-1H,5H pirido[3,2,1-*l*] quinoline).

CLAIMS

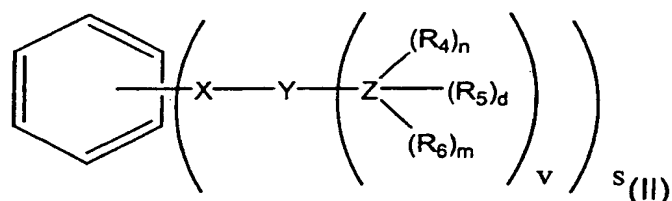
1. Compounds of general formula (I)



5

wherein

R is the following group of formula (II)



10 wherein

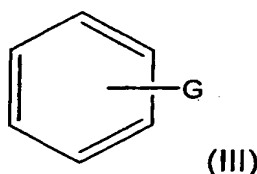
X is selected from the group consisting of O, S, CH₂, COO, CH₂CO, O(CH₂)₂O, O(CH₂)₃O and N;

Z is selected from between N and CH₂N;

Y is selected from aliphatic groups, linear or branched, saturated or unsaturated, having from 1 to 10 carbon atoms, and phenyl, possibly substituted, or Y forms with Z a saturated or unsaturated heterocycle, possibly substituted, comprising up to two heteroatoms selected from the group consisting of N, O and S;

R₄ and R₅, equal or different from each other, are selected from H and alkyl groups having from 1 to 3 carbon atoms, or they form with the Z group a saturated or unsaturated heterocycle, possibly substituted, comprising up to two heteroatoms selected from the group consisting of N, O and S;

- R_6 is selected from H and aliphatic groups, linear or branched, saturated or unsaturated, having from 1 to 5 carbon atoms, possibly substituted with alkylamine or alkylammonium groups having alkyl chains comprising from 1 to 5 carbon atoms, or forming a saturated heterocycle comprising up to two heteroatoms
- 5 selected from between O and N;
- d, m, and n, equal or different from each other, are selected from 0 and 1;
- v and s, equal or different from each other, are integers comprised between 1 and 3;
- R_1 is selected from H and a group of formula (III)



10

wherein

G is selected from H and $P-(CH_2)_l-(W)_f-J$, wherein

P is selected from the group consisting of O, CH_2 , CO_2 , $NHCONH$ and $CONH$;

l is an integer comprised between 0 and 5;

- 15 W is selected from the group consisting of O, CO_2 , $CONH$ and $NHCONH$;

f is selected from between 0 and 1;

J is H or an alkyl group $(CH_2)_q-CH_3$, wherein q is an integer comprised between 0 and 20;

R_2 and R_3 , equal or different from each other, are selected from between R and

- 20 R_1 , wherein R and R_1 are defined as above,

M is chosen from 2H and a metal selected from the group consisting of Zn, Mg, Pt, Pd, $Si(OR_7)_2$, $Ge(OR_7)_2$ and $AlOR_7$, wherein R_7 is chosen from between H and C1-C15 alkyl,

and pharmaceutically acceptable salts thereof,

- 25 with the exception of the following compounds:

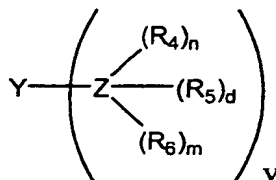
a) compound of formula (I) wherein M is 2H, $R_1 = R_3 = H$, $R = R_2$ is a group of formula (II) in which s is 1, X is O, Y is $(CH_2)_3$, v is 1, Z is N, $n = d = 1$, m is 0, and $R_4 = R_5 = H$; and

b) compound of formula (I) wherein M is 2H, $R_1 = R_3 = H$, $R = R_2$ is a group of formula (II) in which s is 1, X is O, Y is $(CH_2)_3$, v is 1, Z is N, $n = d = 1$, m is 0, R_4 and R_5 form with Z a phthalimido group.

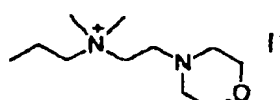
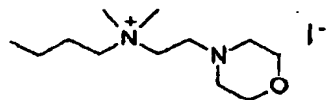
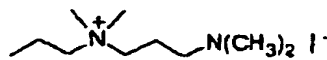
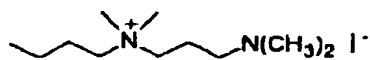
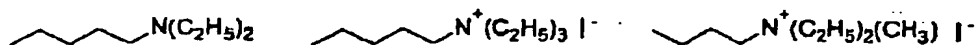
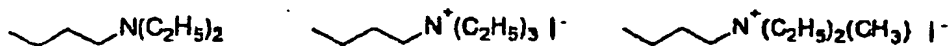
2. Compounds of general formula (I) according to claim 1, in which the said group
 5 R comprises at least one substituent bearing tertiary or quaternary nitrogen.

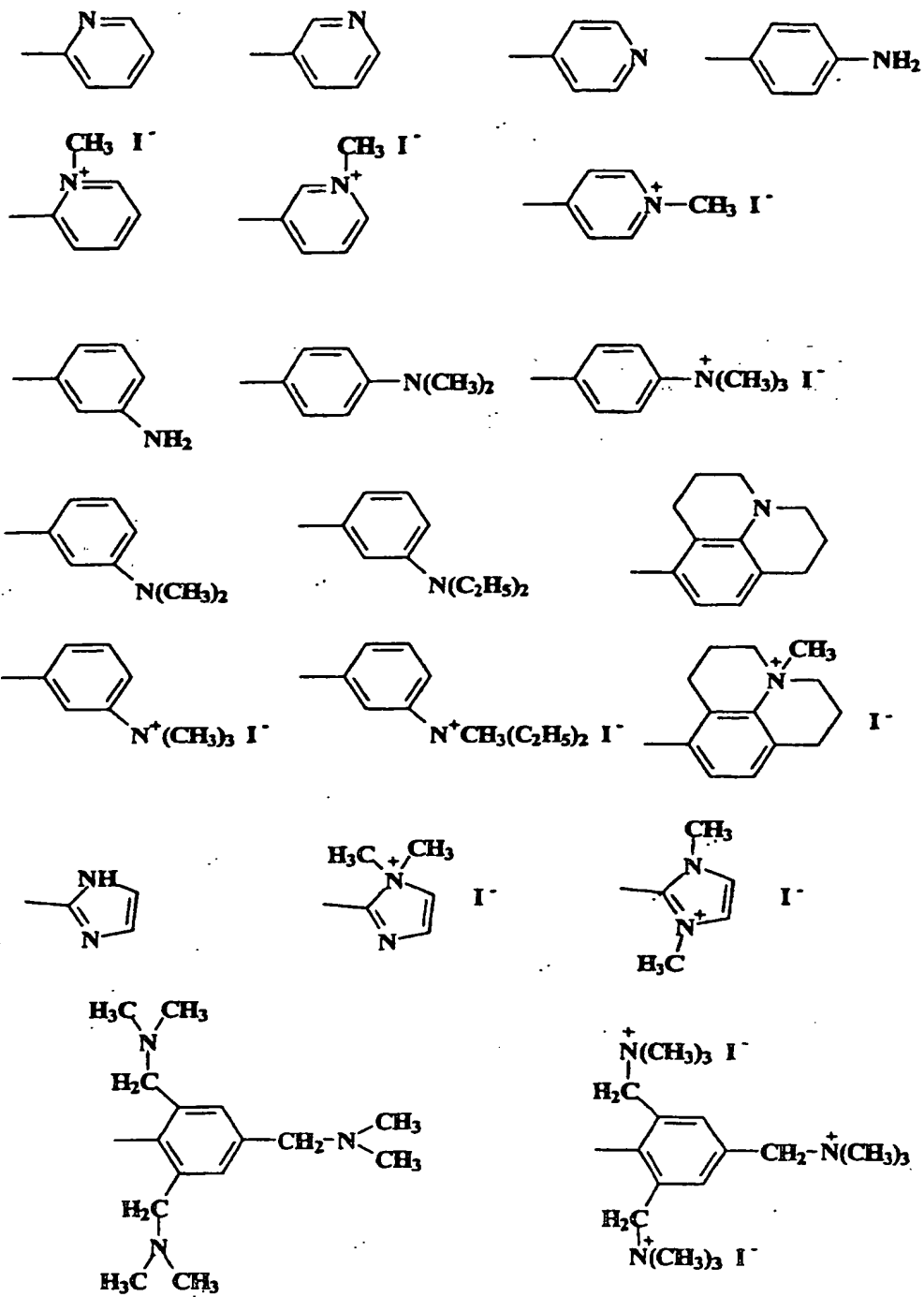
3. Compounds of general formula (I) according to claim 1, wherein said saturated
 or unsaturated heterocycle, possibly substituted, is selected from the group
 consisting of morpholine, piperidine, pyridine, pyrimidine, piperazine, pyrrolidine,
 pyrroline, imidazole, aniline and julolidine (2,3,6,7-tetrahydro-1H,5H-pirido[3,2,1-*lj*]
 10 quinoline).

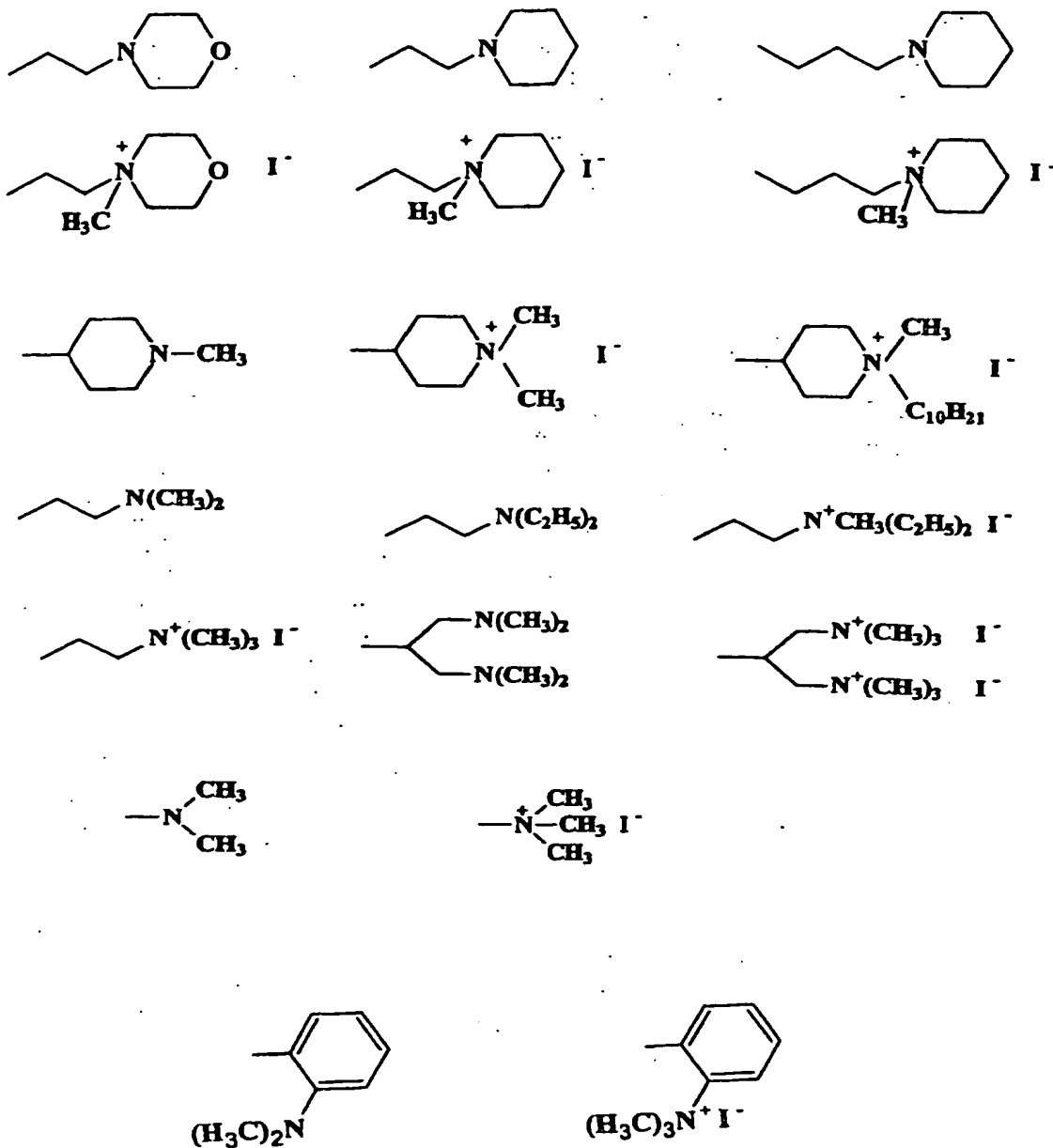
4. Compounds of general formula (I) according to claim 1, wherein the group



is selected from the group consisting of:







5. Compounds of general formula (I) according to claim 1, selected from the group consisting of:

5,10,15-tris-[4-(2-N,N,N-trimethylammoniummethoxy)-phenyl]-20-[(4-decyloxy)-phenyl] porphyrin triiodide,

5 5,10,15-tris-[4-(2-N,N,N-trimethylammoniummethoxy)-phenyl]-20-[(4-decyloxy)-phenyl]porphyrinate zinc (II) triiodide,

5,10,15-tris-[4-(2-N,N-dimethylaminoethoxy)phenyl]-20-[(4-decyloxy)phenyl] porphyrin],

10 5,10,15-tris-[4-(2-N,N-dimethylaminoethoxy)-phenyl]-20-[(4-decyloxy)phenyl] porphyrinate zinc (II),

5,10,15-tris-[[4-(N-methylpiperidin-4-yl)oxy]phenyl]-20-[(4-decyloxy)phenyl] porphyrin,

5,10,15-tris-[[4-(N,N-dimethylpiperidin-4-ium)oxy]phenyl]-20-[(4-decyloxy)phenyl] porphyrin triiodide,

15 5,10,15-tris-[3-(2-morpholin-4-ylethoxy)phenyl]-20-[(4-decyloxy)phenyl]porphyrin,
5,10,15-tris-[[3-(2-methylmorpholin-4-ium)ethoxy]phenyl]-20-[(4-decyloxy)phenyl] porphyrin triiodide,

5,10,15-tris-{4-[4-(N,N-dimethylamino)phenoxy]phenyl}-20-[(4-decyloxy)phenyl] porphyrin,

20 5,10,15-tris-{4-[4-(N,N,N-trimethylammonium)phenoxy]phenyl}-20-[(4-decyloxy)phenyl] porphyrin triiodide,

5,10,15-tris-{4-[3-(N,N-dimethylamino)phenyl]thiophenyl}-20-[(3-undecyloxy) phenyl] porphyrin,

25 5,10,15-tris-{4-[3-(N,N,N-trimethylammonium)phenyl]thiophenyl}-20-[(4-undecyloxy) phenyl]porphyrin triiodide,

5,10,15-tris-[3-(3-N,N-dimethylaminopropoxy)phenyl]-20-[(3-undecyloxy) phenyl] porphyrin,

5,10,15-tris-[3-(3-N,N,N-trimethylammoniumpropoxy)phenyl]-20-[(3-undecyloxy) phenyl] porphyrin triiodide,

30 5,10,15-tris-{4-[4-(N,N-dimethylamino)butoxy]phenyl}-20-[(4-undecyloxy) phenyl] porphyrin,

- 5,10,15-tris-{4-[4-(N,N,N-trimethylammonium)butoxy]phenyl}-20-[(4-decyloxy)phenyl]porphyrin triiodide,
- 5-{4-[2,4,6-tris-[(dimethylamino)methyl]phenoxy]phenyl}-10,15,20-tris-[(4-decyloxy)phenyl]porphyrin,
- 5-{4-[2,4,6-tris-[(trimethylammonium)methyl]phenoxy]phenyl}-10,15,20-tris-[(4-decyloxy)phenyl]porphyrin triiodide,
- 5-{3-[2-(dimethylamino)]-1-[(dimethylamino)methyl]ethoxy}phenyl}-10,15,20-tris-[(3-decyloxy)phenyl]porphyrin,
- 5-{3-[2-(trimethylammonium)]-1-[(trimethylammonium)methyl]ethoxy}phenyl}-10,15,20-tris-[(3-decyloxy)phenyl]porphyrin diiodide,
- 5,10,15-tris-{4-[3-(diethylamino)propoxy]phenyl}-20-[(4-decyloxy)phenyl]porphyrin,
- 5,10,15-tris-{4-[3-(trimethylammonium)propoxy]phenyl}-20-[(4-decyloxy)phenyl]porphyrin triiodide,
- 5,10,15-tris-[4-(2-aminoethoxy)phenyl]-20-[(4-decyloxy)phenyl]porphyrin,
- 5,10,15-tris-{[4-(2-trimethylammonium)ethoxy]phenyl}-20-[(4-decyloxy)phenyl]porphyrin triiodide,
- 5,10,15-tris-{[4-(N,N,N-trimethylammonium)phenoxy]carbonyl}phenyl}-20-[(4-decyloxy)phenyl]porphyrin triiodide,
- 5-{4-{2-(trimethylammonium)-1-[(trimethylammonium)methyl]ethoxy}carbonyl}phenyl}-10,15,20-tris-[(3-decyloxy)phenyl]porphyrin diiodide,
- 5,15-bis-[3-(3-N,N,N-trimethylammoniumpropoxy)phenyl]porphyrin diiodide,
- 5,15-bis-[4-(2-piperidin-1-ylethoxy)phenyl]porphyrin,
- 5,15-bis-[4-(2-N-methylpiperidin-1-iumethoxy)phenyl]porphyrin diiodide,
- 5,15-bis-[4-(3-N,N-dimethylaminopropoxy)phenyl]-10,20-bis-[(3-decyloxy)phenyl]porphyrin,
- 5,15-bis-[4-[3-N,N,N-trimethylammoniumpropoxy]phenyl]-10,20-bis-[(3-decyloxy)phenyl]porphyrin diiodide,
- 5,15-bis-4-{[2-(N,N-dimethylamino)ethylthio]phenyl}porphyrin,
- 5,15-bis-[4-[2-(N,N,N-trimethylammonium)ethylthio]phenyl]porphyrin diiodide,
- 5,15-bis-[4-[2-[3-(trimethylammonium)phenoxy]ethoxy]phenyl]porphyrin diiodide,
- 5,15-bis-[4-[2-[3-(N,N,N-trimethylammonium)phenyl]-2-oxoethyl]-10,20-bis-[(3-decyloxy)phenyl]porphyrin diiodide,

5,15-bis-[3-(3-N,N,N-trimethylammoniumpropoxy)phenyl]porphyrinate zinc(II) diiodide,

5,15-bis-[3-(3-N,N-dimethylaminopropoxy)phenyl]porphyrinate zinc(II),

5,15-bis-[4-(4-N,N,N-trimethylammoniumphenoxy)phenyl] porphyrin diiodide,

5 5,15-bis-[4-(4-aminophenoxy)phenyl]porphyrin,

5,15-bis-[3-(4-N,N-dimethylaminophenoxy)phenyl]porphyrin,

5,15-bis-[3-(4-N,N,N-trimethylammoniumphenoxy)phenyl]porphyrin diiodide,

5,15-bis-[3-(4-N,N-dimethylaminophenyl)thiophenyl]porphyrin,

5,15-bis-[3-(4-N,N,N-trimethylammoniumthiophenoxy)phenyl]porphyrin diiodide,

10 5,15-bis-4-[3-(N,N-dimethylaminophenoxy)phenyl]-10,20-bis-[(4-decyloxy)phenyl]porphyrin,

5,15-bis-4-[3-(N,N,N-trimethylammoniumphenoxy)phenyl]-10,20-bis-[(4-decyloxy)phenyl]porphyrin diiodide,

15 5,10,15-tris-{4-[4-(N,N-dimethylamino)butoxy]phenyl}-20-[(4-undecyloxy)phenyl]porphyrinate zinc(II),

5,10,15-tris-{4-[4-(N,N,N-trimethylammonium)butoxy]phenyl}-20-[(4-undecyloxy)phenyl]porphyrinate zinc(II) triiodide,

5,15-bis-[4-(2-piperidin-1-ylethoxy)phenyl]porphyrinate zinc(II), and

5,15-bis-[4-(2-N-methylpiperidin-1-iumethoxy)phenyl]porphyrinate zinc(II) diiodide.

20 6. Conjugates of compounds of general formula (I) as defined in claims 1-5 with a macromolecule selected from the group consisting of aminoacids, polypeptides, proteins and polysaccharides.

7. Process for the preparation of compounds of formula (I) in which $R = R_2 = R_3$ as defined in claims 1-5, selected from the group consisting of:

25 - pre-functionalization of suitable reagents with amino groups, followed by statistical synthesis of the porphyrin ring, possible modification of the amino groups in ammonium groups, and possible complexation with the metal cation if the metal complex is required;

30 - statistical synthesis with formation of the porphyrin ring followed by functionalization of the porphyrin with the present amino or ammonium groups, and possible complexation with the metal cation; and

- synthesis of the porphyrin ring through suitable dipyrromethane derivatives followed by functionalisation of the porphyrin with the present amino or ammonium groups, and possible complexation with the metal cation.

8. Process for the preparation of compounds of formula (I) in which $R = R_2$ and $R_1 = R_3$ as defined in claims 1-5, comprising the synthesis of the porphyrin ring through dipyrromethane followed by functionalisation of the porphyrin with aliphatic or aromatic amino or ammonium groups, and possible complexation with the metal cation if the metal complex is required.

9. Intermediate compounds in the preparation of compounds of formula (I) as defined in claims 1-5, selected from the group consisting of:

5,10,15-tris-[4-(2-hydroxyethoxy)phenyl]-20-[(4-decyloxy)phenyl]porphyrin,
5,10,15-tris-[4-(2-methylsulphonylethoxy)phenyl]-20-[(4-decyloxy)phenyl]porphyrin,
5,15-bis-[3-(3-hydroxypropoxy)phenyl]porphyrin,
5,15-bis-[3-(3-methylsulphonylpropoxy)phenyl]porphyrin,
5,15-bis-[3-(3-hydroxypropoxy)phenyl]porphyrinate zinc(II),
5,15-bis-[3-(3-methylsulphonylpropoxy)phenyl]porphyrinate zinc(II),
5,15-bis-[[3-(3-(4-methylphenyl)sulfonyl)oxy]propoxyphenyl]porphyrinate zinc(II),
5,15-bis-[3-(3-bromopropoxy)phenyl]porphyrinate zinc(II), and
5,15-bis-[4-(4-nitrophenoxy)phenyl]porphyrin.

10. Pharmaceutical compositions comprising as the active principle at least a compound of general formula (I) as defined in claims 1-5, or a conjugate according to claim 6, or mixtures thereof, possibly in combination with pharmaceutically acceptable excipients and/or diluents.

11. Use of compounds of general formula (I) as defined in claims 1-5, or of conjugates thereof according to claim 6, for the preparation of pharmaceutical compositions for photodynamic therapy.

12. Use of compounds of general formula (I) or of conjugates thereof according to claim 11, for the preparation of pharmaceutical compositions for the treatment of bacterial, viral or micotic infections.

13. Use of compounds of general formula (I) or of conjugates thereof according to claim 11, for the preparation of pharmaceutical compositions for the treatment of diseases characterised by cellular hyperproliferation.

14. Use of compounds of general formula (I) according to claim 13, wherein said diseases characterised by cellular hyperproliferation are selected from the group consisting of psoriasis, intimal hyperplasia, benign prostate hyperplasia and atheromas.

5 15. Diagnostic agents comprising as the active principle a compound of general formula (I) as defined in claims 1-5, or a conjugate thereof according to claim 6, possibly in combination with a pharmaceutically acceptable carrier.

16. Sterilizing agents comprising as the active principle a compound of general formula (I) as defined in claims 1-5, or a conjugate thereof according to claim 6,
10 possibly in combination with a pharmaceutically acceptable carrier.

17. Use of a compound of general formula (I) as defined in claims 1-5, or a conjugate thereof according to claim 6, for blood and blood derivatives sterilisation.

18. Use of a compound of general formula (I) as defined in claims 1-5, or a
15 conjugate thereof according to claim 6, for the preparation of a pharmaceutical composition for the sterilisation of wounds.

19. Method of treating infectious diseases of viral, fungine and bacterial origin, diseases characterised by cellular hyperproliferation and dermatological diseases, comprising administering to a patient in need of such a treatment an effective
20 amount of at least a compound of general formula (I) as defined in claim 1 or a conjugate thereof according to claim 6, and irradiating the pathologically affected tissues with light of appropriate wavelength.

20. Method according to claim 19, wherein the said affected tissues are irradiated by visible red light radiation when the treatment of deep seated tumours on
25 infections is required, and by blue visible radiation or white light radiation when treating psoriasis, actinic keratoses, basal cell carcinomas and other cancerous and pre-cancerous lesions of the skin and mucosas.

21. Method of localising pathologically affected areas comprising administering to a patient an effective amount of at least a compound of general formula (I) as
30 defined in claim 1 or a conjugate thereof according to claim 6, and irradiating the pathologically affected areas with light of appropriate wavelength.